

Scientists identify gene as master regulator in schizophrenia

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An illustration of the study design to identify and validate TCF4 as a key gene in schizophrenia. Credit: Abolfazl Doostparast Torshizi, Jubao Duan and Kai Wang

Using computational tools to investigate gene transcription networks in large collections of brain tissues, a scientific team has identified a gene that acts as a master regulator of schizophrenia during early human brain development. The findings may lay the groundwork for future



treatments for the highly complex neuropsychiatric disorder.

"Because hundreds, or even thousands, of <u>genes</u> may contribute to the risk of schizophrenia, it is crucial to understand which are the most important ones, orchestrating core networks in the disease," said study leader Kai Wang, Ph.D., of the Department of Pathology and Laboratory Medicine and the Raymond G. Perelman Center for Cellular and Molecular Therapeutics (CCMT) at Children's Hospital of Philadelphia (CHOP). "Pinpointing master regulators may help guide us toward priority targets for novel treatments in the future."

The study appeared online today in *Science Advances*. Wang co-led the research with Jubao Duan, Ph.D., the Charles R. Walgreen Research Chair and an associate professor at the Center for Psychiatric Genetics of North Shore University HealthSystem (NorthShore) and the University of Chicago, Evanston, Ill. The first author was Abolfazi Doostparast Torshizi, Ph.D., of the CCMT at CHOP.

Although schizophrenia affects about 1 in 100 adults, and is highly heritable, the genetic architecture of the neuropsychiatric disorder is notoriously complex, with many unsolved questions. Wang and colleagues contrast the current state of knowledge to recent advances in cancer research. Oncology researchers have identified many gene pathways and networks that when perturbed, result in specific types of cancer. That knowledge has permitted fine distinctions in diagnosing subsets among types of cancer, and has led to individualized treatments based on a patient's genetic profile.

The underlying genomic biology in neuropsychiatric disorders is likely more challenging. In the most extreme case, researchers have recently proposed an "omnigenic" model, in which nearly all the genes in a disease-relevant cell type contribute to a specific neuropsychiatric disorder, but Wang points out, "Not all the genes carry equal



weight—the problem is to determine which are more important than others."

Thus, the study team used computational systems biology approaches to discern a disease-relevant core pathway in schizophrenia and to discover a master regulator in that pathway that affects hundreds of downstream genes.

Wang and colleagues analyzed two independent datasets of biological samples from schizophrenia patients and control subjects. One dataset, the CommonMind Consortium (CMC), is a public-private partnership with well-curated brain collections. The other was a collection of primary cultured neuronal cells derived from olfactory epithelium (CNON), generated by study co-authors at the University of Southern California and SUNY Downstate. The CMC dataset contained adult postmortem brain tissue, while the CNON dataset, used to validate findings from the CMC study, represented <u>cell cultures</u> that contain <u>neuronal cells</u> from nasal biopsies. Applying an algorithm developed at Columbia University to reconstruct gene transcription networks, the study team identified the gene *TCF4* as a master regulator for schizophrenia.

Previous genome-wide association studies (GWAS) had indicated that *TCF4* was a locus for schizophrenia risk, said Wang, but little was known of the gene's functional effects. The study team investigated those effects by knocking down, or decreasing, the gene's expression in neural progenitor cells and glutamatergic neurons derived from induced pluripotent stem cells in Duan's lab at NorthShore.

Observations on three different cell lines showed that, when knocked down, the predicted *TCF4* regulatory networks were enriched for genes showing transcriptomic changes, as well as for genes involved in neuronal activity, schizophrenia risk genes having genome-wide



significance, and schizophrenia-associated de novo mutations. Although some of the cellular effects of *TCF4* dysregulation were previously shown in mice, Duan noted that the results from perturbing *TCF4* gene networks in human stem cell models may be more relevant to the neurodevelopmental aspects of neuropsychiatric disorders.

The current study, said the researchers, sets the stage for further investigations. One direction, said Wang, is to use expanded datasets to explore whether other master regulators in addition to *TCF4* may act in schizophrenia. If so, it may eventually be possible to classify patients with schizophrenia into subgroups more responsive to specific treatments, as is occurring in many cancers, to assist in implementing precision medicine in psychiatric diseases.

Other approaches, added Doostparast, may involve pursuing functional genomics at the level of single <u>cells</u>, to assess the cell types that are most influenced by dysregulating gene expression.

The study represents one of the first successful examples of combining computational approaches and stem cell-based experimental models to disentangle complex gene networks in psychiatric diseases.

Wang noted limitations in the study. The empirical validation focused on <u>neural progenitor cells</u> and glutamatergic neurons, but other cell types may be investigated in the future, such as interneurons and microglia, which have also been implicated in the development of <u>schizophrenia</u>.

More information: A. Doostparast Torshizi el al., "Deconvolution of transcriptional networks identifies TCF4 as a master regulator in schizophrenia," *Science Advances* (2019). <u>DOI: 10.1126/sciadv.aau4139</u>, <u>advances.sciencemag.org/content/5/9/eaau4139</u>



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